L	Hits	Search Text	DB	Time stamp
Number	5.			
1	0	dinh-tan-thanh-\$.in.	USPAT;	2002/07/31
			US-PGPUB	09:43
2	2	dinh-tan-thanhin.	USPAT;	2002/07/31
_			US-PGPUB	09:43
3	0	tremble-patrice-\$.in.	USPAT;	2002/07/31
			US-PGPUB	09:44
4	1	tremble-patricein.	USPAT;	2002/07/31
3	_	Cichare passion van	US-PGPUB	09:44
5	11	cunanan-crystal-\$.in.	USPAT;	2002/07/31
3	11	Cultalian Clystal V.III.	US-PGPUB	09:44
6	0	cunanan-crystal-m-\$.in.	USPAT;	2002/07/31
6	٥	Cullanan-Crystar-m-7.111.	US-PGPUB	09:44
_			USPAT;	2002/07/31
7	11	cunanan-crystal-min.	US-PGPUB	09:44
_			USPAT;	2002/07/31
8	0	may-christine-\$.in.	1	09:44
	_		US-PGPUB	2002/07/31
9	0	may-christinein.	USPAT;	
,			US-PGPUB	09:44
10	0	((phospholipid\$1) and (chromatagraphy or	USPAT;	2002/07/31
		tlc)).ti.	US-PGPUB	09:45
11	. 260	(phospholipid\$1) same (chromatagraphy or	USPAT;	2002/07/31
		tlc)	US-PGPUB	09:46
12	107	((phospholipid\$1) same (chromatagraphy or	USPAT;	2002/07/31
		tlc)) same (thin adj layer)	US-PGPUB	09:46
13	6	(((phospholipid\$1) same (chromatagraphy	USPAT;	2002/07/31
		or tlc)) same (thin adj layer)) same	US-PGPUB	09:50
		((one adj dimensional) or (one adj way)		
		or (one adj direction))		
14	10	(((phospholipid\$1) same (chromatagraphy	USPAT;	2002/07/31
		or tlc)) same (thin adj layer)) and	US-PGPUB	09:53
		436/71.ccls.		
15	147759	((acetic adj acid) or ch3cooh)	USPAT;	2002/07/31
13	14/173	((access auj actu, of chocodi,	US-PGPUB	09:54
16	1613	(((acetic adj acid) or ch3cooh)) same	USPAT;	2002/07/31
10	1013	(((acetic ad) acid) of chiscoon, same ((potassium adj chloride) or kcl)	US-PGPUB	09:54
1.7	60	((((acetic adj acid) or ch3cooh)) same	USPAT;	2002/07/31
17	00	(((lacetic adj acid) of (inscoon)) same	US-PGPUB	09:54
		((potassium adj chloride) or kcl)) same	03-1910	09.53
		(chromatography or tlc)	HCDAM.	2002/07/31
18	14	(((((acetic adj acid) or ch3cooh)) same	USPAT;	
		((potassium adj chloride) or kcl)) same	US-PGPUB	09:57
		(chromatography or tlc)) same (solvent\$1		
ľ		or (mobile adj phase))		

WEST Search History

DATE: Wednesday, July 31, 2002

Set Name	Hit Count	Set Name result set			
DB=JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=ADJ					
L17	L16 and chromatography	7	L17		
L16	L15 and ((potassium adj chloride) or kcl)	219	L16		
L15	(acetic adj acid) or ch3cooh	46629	L15		
L14	L13 and ((one adj dimensional) or (one adj way) or (one adj direction))	0	L14		
L13	phospholipid\$1 and (tlc or ((thin adj layer) adj chromatography))	39	L13		
L12	wo-9950655-\$.did.	1	L12		
LII	wo-9950655-\$.in.	0	L11		
L10	wo-0233399-\$.did.	0	L10		
L9	wo-02033399-\$.did.	0	L9		
L8	wo-2002033399-\$.did.	0	L8		
L7	L6 and phospholipid\$1	0	L7		
L6	may-c-\$.in.	148	L6		
L5	L4 and phospholipid\$1	0	L5		
L4	cunanan-c-\$.in.	13	L4		
L3	tremble-p-\$.in.	0	L3		
L2	L1 and phospholipid\$1	0	L2		
L1	dinh-t-\$.in.	24	Ll		

END OF SEARCH HISTORY

		E DINH TAN THANH/AU
L1	10	S E1-E3
L2	2	S L1 AND PHOSPHOLIPID?
		E TREMBLE PATRICE/AU
L3	76	S E1-E5
L4	1	S L3 AND PHOSPHOLIPID?
		E CUNANAN CRYSTAL M/AU
L5	16	S E2-E3
L6	2	S L5 AND PHOSPHOLIPID?
		E MAY CHRISTINE/AU
L7	1	S E4
F8	2975	S PHOSPHOLIPID? AND (THIN LAYER CHROMATOGRAPY OR TLC)
L9	56	S L8 AND (ONE DIMENSIONAL OR ONE WAY OR ONE DIRECTION)
L10	5	S L9 AND (POTASSIUM CHLORIDE OR KCL)
L11		S L10 AND ACETIC ACID
L12		S L10 NOT L11
L13		S L9 AND PRIMULIN
L14	3	DUP REMOV L13 (2 DUPLICATES REMOVED)
L15	51	S L9 NOT L10
L16		S L15 NOT L13
L17		DUP REMOV L16 (8 DUPLICATES REMOVED)
L18		S ACETIC ACID OR CH3COOH
L19		S L18 AND (POTASSIUM CHLORIDE OR KCL)
L20		S L19 AND (CHROMATOGRAPHY OR TLC)
L21		S L20 AND (SOLVENT? OR MOBILE PHASE?)
L22	12	DUP REMOV L21 (2 DUPLICATES REMOVED)

lysophosphatidylcholine) and three lysophospholipids (lysophosphatidylserine, lysophosphatidylethanolamine and lysophosphatidylcholine). This is achieved by simple involvement of 0.4% ammonium sulfate in silica gel H and of acetone in a developing solvent as chloroform-methanol-acetic acid-acetone-water (40:25:7:4:2). The procedure is simple and the sepn. is reproducible. The weakness of this method is the partial degrdn. of phosphatidylethanolamine to lysophosphatidylethanolamine, but a method to prevent this degrdn. is also presented.

L12 ANSWER 8 OF 12 CAPLUS COPYRIGHT 2002 ACS

Full elates Text Palakances

AN 1992:3013 CAPLUS

DN 116:3013

TI Comparison of mobile phases for separation of **phospholipids** by **one-dimensional TLC** on preadsorbent high performance silica gel plates

AU Aloisi, Jacqueline; Fried, Bernard; Sherma, Joseph CS Dep. Biol., Lafayette Coll., Easton, PA, 18042, USA

SO J. Liq. Chromatogr. (1991), 14(18), 3269-75

CODEN: JLCHD8; ISSN: 0148-3919

DT Journal

LA English

Eight solvent systems reported in the literature for the 1-dimensional TLC sepn. of phospholipids were compared under identical conditions by using high-performance preadsorbent silica gel plates. The best overall sepn. of phospholipid stds. was obtained by a single development with chloroform-methanol-water (65:25:4), and 3 other systems contg. chloroform also gave good sepns. Rf Data are tabulated for these 4 systems, and the phospholipids extd. from the digestive gland-gonad complex of Biomphalaria glabrata snails are identified.

L12 ANSWER 9 OF 12 CAPLUS COPYRIGHT 2002 ACS

Full clang Text References

AN 1991:181515 CAPLUS

DN 114:181515

TI Comparison of mobile phases for separation and quantification of lipids by one-dimensional TLC on preadsorbent high performance silica gel plates

AU Aloisi, Jacqueline D.; Sherma, Joseph; Fried, Bernard

CS Dep. Biol., Lafayette Coll., Easton, PA, 18042, USA

SO J. Liq. Chromatogr. (1990), 13(20), 3949-61 CODEN: JLCHD8; ISSN: 0148-3919

DT Journal

LA English

Twenty-four solvent systems reported in the literature for the 1-dimensional TLC sepn. of lipids and phospholipids were compared under identical conditions by using high-performance preadsorbent silica gel plates. The best overall sepn. of mixts. of neutral lipid and phospholipid stds. and compds. extd. from the digestive gland-gonad complex of Biomphalaria glabrata snails was obtained with a system utilizing consecutive development with CHCl3-MeOH-H2O (65:25:4), CHCl3-hexane (3:1), and CCl4. The best system for quantification of neutral lipids was hexane-Et2O-HCOOH (80:20:2). Rf Data are tabulated and results discussed for all systems tested.

L12 ANSWER 10 OF 12 MEDLINE



AN 83111030 MEDLINE

DN 83111030 PubMed ID: 6822837

TI Receptor-mediated increases in phosphatidylinositol turnover in neuron-like cell lines.

L10 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2002 ACS

EU (Chainsin References Text

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2002:315200 CAPLUS
AN
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DN 136:291337

Methods for quantitative and qualitative analyses of phospholipids using ΤI one-dimensional thin layer chromatography

Dinh, Tan Thanh; Tremble, Patrice; Cunanan, Crystal M.; Cabiling, IN Christine May

Edwards Lifesciences Corporation, USA PA

PCT Int. Appl., 23 pp. SO

CODEN: PIXXD2

Patent DT

English LA

FAN.CNT 1

APPLICATION NO. KIND DATE PATENT NO. ------_____ ____ WO 2001-US32023 20011012 20020425 A2 WO 2002033399 ΡI W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG PRAI US 2000-693186 20001019 Α

A highly sensitive and specific method for the detection and quantification of lipids is provided. Specifically, methods for the simultaneous detection and quantification of phospholipids extd. from mammalian tissues is described. The anal. methods provided disclose a modified one-dimensional thin-layer chromatog. technique specifically developed to rapidly and accurately detect and quantify phospholipids from mammalian cardiac tissues.

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2002 ACS

Full Feferences Text

1991:181515 CAPLUS AN

114:181515

DN Comparison of mobile phases for separation and quantification of lipids by one-dimensional TLC on preadsorbent high performance silica gel plates

Aloisi, Jacqueline D.; Sherma, Joseph; Fried, Bernard ΑU

Dep. Biol., Lafayette Coll., Easton, PA, 18042, USA CS

J. Liq. Chromatogr. (1990), 13(20), 3949-61 SO CODEN: JLCHD8; ISSN: 0148-3919

DTJournal

English LA

Twenty-four solvent systems reported in the literature for the AB 1-dimensional TLC sepn. of lipids and phospholipids were compared under identical conditions by using high-performance preadsorbent silica gel plates. The best overall sepn. of mixts. of neutral lipid and phospholipid stds. and compds. extd. from the digestive gland-gonad complex of Biomphalaria glabrata snails was obtained with a system utilizing consecutive development with CHCl3-MeOH-H2O (65:25:4), CHCl3-hexane (3:1), and CCl4. The best system for quantification of neutral lipids was hexane-Et20-HCOOH (80:20:2). Rf Data are tabulated and results discussed for all systems tested.

=>

dimensional thin-layer chromatography

- ΑU White, Thayer; Bursten, Stuart; Federighi, David; Lewis, Robert A.; Nudelman, Edward
- CS Cell Therapeutics, Inc., Seattle, WA, 98119, USA
- Analytical Biochemistry (1998), 258(1), 109-117 SO CODEN: ANBCA2; ISSN: 0003-2697
- PB Academic Press
- DT Journal
- LΑ
- English AΒ An improvement of current methods is needed for simple, rapid, and precise quantification of cellular lipids, including rare species of biol. active cellular lipids, such as phosphatidic acid (PA) and diradylglycerol (DG). In addn., further anal. of hydrolyzed acyl chains from these species by methods such as gas chromatog. requires complete sepns. Methods have been developed for the quantification of neutral lipids and several phospholipids extd. from mammalian cells and sera. Lipid masses were detd. for the major classes of the neutral, nonpolar lipids, and of the phospholipids. The lipid classes were sepd. by a multistep thin-layer chromatog. (TLC) procedure in different solvent systems, a method which we have designated as multi-one-dimensional thin-layer chromatog. (MOD-TLC). Resolved lipid bands were visualized by the lipophilic dye primulin (direct yellow 59) and scanned by an automated laser-fluorescence detector. The mass of each band was detd. by comparing band intensities of unknown samples to diln. curves of authentic stds. With modifications in solvent mixts. and length of sepn. times, the majority of biol. lipids could be resolved and quantified with MOD-TLC methods. Since the detection method is nondestructive, purified lipids could then be recovered by scraping the visualized bands and extg. the lipids from the silica. The structural identities of the recovered lipids were confirmed by fast-atom bombardment and electrospray mass spectrometry. Extd. lipids were also hydrolyzed to release acylchains and acyl chain species were detd. in comparison to authentic stds. by gas chromatog. PA and DG levels in ECV.304 cells were found to be 4.6 and 3.3%, resp., of PC levels, with a PA/DG ratio of 1.4, which is in accord with published experience using other methods and different cell types. PA in human serum was detected at 0.6% of PC, indicating the sensitivity of the technique. In contrast to two-dimensional thin-layer chromatog., which allows for good resoln. of some lipid species, but cannot be used to analyze more than a single exptl. point per plate, MOD-TLC allows for direct comparative anal. of multiple samples on a single TLC plate, while still providing good resoln. for the quantification of most major classes of lipid species.

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UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2000-693186
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                            20001019
     A highly sensitive and specific method for the detection and
     quantification of lipids is provided. Specifically, methods for the
     simultaneous detection and quantification of phospholipids extd. from
     mammalian tissues is described. The anal. methods provided disclose a
     modified one-dimensional thin-layer chromatog. technique specifically
     developed to rapidly and accurately detect and quantify phospholipids
     from mammalian cardiac tissues.
L14 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2002 ACS
         Peferences
   Text
     1999:641071 CAPLUS
ΑN
DN
     131:269265
    Methods of separation and detection of hydrophobic target molecules by
    multiple one-dimensional thin layer chromatography
    White, Thayer; Nudelman, Edward D.
·ΙΝ
    Cell Therapeutics, Inc., USA
PΑ
     PCT Int. Appl., 41.pp.
SO
     CODEN: PIXXD2
DT
     Patent
LΑ
    English
FAN.CNT 2
     PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
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    WO 9950655
                                          WO 1999-US6803
                                                           19990330
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         W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
             DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
             JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
             MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
             TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ,
             MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK,
             ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,
             CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     AU 9934548
                            19991018
                                            AU 1999-34548
                       A1
                                                              19990330
US 6331254
PRAI US 1998-49941
                             20011218
                                            US 1999-465678
                       В1
                                                              19991217
                       A1
                             19980330
     WO 1999-US6803
                       W
                             19990330
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AB Methods which employ thin layer chromatog. for sepg. and detecting hydrophobic target mols. are particularly useful in sepg. biol. relevant lipids. By utilizing non-destructive detection techniques, these methods also can be adapted to further quantification or structural anal. Lipids extd. from ECV.304 cells and from pooled human serum samples were sepd. by multiple one-dimensional (MOD) TLC sequentially using chloroform-methanol-acetic acid (90:10:1, vol./vol./v), hexane-diethylether-acetone (60:40:5, vol./vol./v), hexane-diethylether (97:3, vol./vol.), and hexane (100%) as the mobile phases, all run in the same direction. The dried plates were sprayed with Primulin dye soln. and scanned by laser-excited fluorescent detection.

RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2002 ACS

DUPLICATE 1



AN 1998:236214 CAPLUS

DN 129:2268

TI High-resolution separation and quantification of neutral lipid and phospholipid species in mammalian cells and sera by multi-one-